



California Drug Recall Information



Recall Name

**Hospira Recalls One Lot Of Labetalol Hydrochloride Injection, USP,
100 MG/20 ML (5MG/ML), 20 ML, Multidose Vials
Due to Visible Particulates**

Recall Date	Product Description	Recalling Firm	Recall Reason
05/16/14	Labetalol Hydrochloride Injection, USP 100 MG/20 ML (5MG/ML) 20 ML Multidose Vials NDC 0409-2267-20	Hospira, Inc. Lake Forest, IL	<i>Confirmed customer report of embedded particulate within the glass vial and visible particles floating in the solution.</i>
Recall Class	Product Identification	Distribution	Affected Dates
N/A	Suspect Lot: <ul style="list-style-type: none">Lot 36-225-DD Expiration: 12/01/2015 Product Photo	CA , nationwide	Distributed: February 2014

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/Safety/Recalls/ucm397726.htm>